



Core-shell structured reduced graphene oxide wrapped magnetically separable rGO@CuZnO@Fe₃O₄ microspheres as superior photocatalyst for CO₂ reduction under visible light

Pawan Kumar^a, Chetan Joshi^a, Alexandre Barras^b, Brigitte Sieber^c, Ahmed Addad^c, Luc Boussekey^d, Sabine Szunerits^b, Rabah Boukherroub^{b,*}, Suman L. Jain^{a,*}

^a Chemical Science Division, CSIR-Indian Institute of Petroleum, Dehradun, 248005, India

^b Univ. Lille, CNRS, Centrale Lille, ISEN, Univ. Valenciennes, UMR 8520 - IEMN, F-59000 Lille, France

^c UMET, UMR CNRS 8207, Université Lille 1, 59655 Villeneuve d'Ascq Cédex, France

^d LASIR, UMR CNRS 8516, Université Lille 1, 59655 Villeneuve d'Ascq Cédex, France

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ABSTRACT

A highly efficient, recyclable and magnetically separable core-shell structured CuZnO@Fe₃O₄ microsphere wrapped with reduced graphene oxide (rGO@CuZnO@Fe₃O₄) photocatalyst has been developed and used for the photoreduction of carbon dioxide with water to produce methanol under visible light irradiation. Owing to the synergistic effect of the components and to the presence of a thin Fe₂O₃ layer on Fe₃O₄, rGO@CuZnO@Fe₃O₄ **4** exhibited higher catalytic activity as compared to the other possible combinations such as CuZnO@Fe₃O₄ **2** and GO@CuZnO@Fe₃O₄ **3** microspheres. The yield of methanol in case of using **2** and **3** as photocatalyst was found to be 858 and 1749 μmol g⁻¹ cat, respectively. However, the yield was increased to 2656 μmol g⁻¹ cat when rGO@CuZnO@Fe₃O₄ **4** was used as photocatalyst under similar experimental conditions. This superior photocatalytic activity of **4** was assumed to be due to the restoration of the sp² hybridized aromatic system in rGO, which facilitated the movement of electrons and resulted in better charge separation. The synthesized heterogeneous photocatalyst could readily be recovered by external magnet and successfully reused for six subsequent cycles without significant loss in the product yield.

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1. Introduction

The rising demand for fossil fuels resulted in emissions of large quantities of carbon dioxide (CO₂) in the atmosphere, which has become a severe environmental issue associated with global warming and climate change [1]. Furthermore, the depletion of hydrocarbon fuels is an inevitable problem which has to be faced in the near future. Therefore, conversion of CO₂ into value-added chemicals such as hydrocarbons is of great importance that can help in tackling both environmental as well as energy issues. More importantly, CO₂ conversion into fuel (energy) that not only mitigates CO₂ emissions, but also recycles it back as a fuel feedstock is highly desirable in meeting clean energy demand [2–4].

CO₂ conversion can be achieved through different means based on chemical, electrochemical, biological and photochemi-

cal approaches. Among them, biomimetic artificial photosynthesis has recently gained considerable attention as a promising technology for reduction of CO₂ to fuel using solar energy [4–6]. In this process, the hydrocarbons are produced by photoreduction of CO₂ using water and solar energy. This method has gained considerable importance in recent years as it is economically viable and environmentally friendly. A large variety of semiconductor materials such as TiO₂, ZnO, ZrO₂, CdS, Fe₂O₃, SiC, WO₃ and transition metal complexes have been extensively investigated for photocatalytic conversion of CO₂ [4,7]. Since redox potentials of large band gap semiconductors are more adapted for CO₂ photoreduction, TiO₂-based photocatalysts have been widely investigated for this transformation [7,8]. Even though higher yields were obtained with metal complexes as compared to TiO₂ photocatalysts, metal complexes, besides being expensive and unstable under UV light, often required an electron donor (e.g. triethylamine), whereas water molecules were used as electron donor for TiO₂ [8]. Band gap engineering by doping with metals and non metals (Cu, Ag, Ru, N, etc.) was used for improvement of the conversion efficiency, but still

* Corresponding authors.

E-mail address: suman@iip.res.in (S.L. Jain).

no satisfactory improvements were achieved [9–12]. Zinc oxide (ZnO), although less investigated than TiO_2 for CO_2 photoreduction, is a wide band gap (3.37 eV) material which can be established as an interesting alternative due to its superior photocatalytic activity, low cost and environmentally friendly properties. In order to extend the operation conditions to higher wavelength, metallic or non metallic elements can be introduced to ZnO material. Yarahmadi and Sharifnia successfully used dye photosensitized ZnO with Co, Ni and Cu phthalocyanines (Pc) for the photocatalytic conversion of CO_2 and CH_4 under visible light [13]. Formation of ZnO/ZnTe cation heterostructure exhibited good photocatalytic activity for the reduction of CO_2 into CH_4 under visible light [14].

Recently, graphene oxide (GO), reduced graphene oxide (rGO) and their nanocomposites have emerged as interesting materials with potential applications in various fields [15–17]. Hsu et al. were the first to report the photocatalytic efficiency of GO for the conversion of CO_2 to methanol under visible light and the conversion rate was found to be 6-fold higher than commonly used TiO_2 photocatalyst [18]. The combination of graphene with semiconductor photocatalysts provided nanocomposites which have shown improved photocatalytic performance for CO_2 photoreduction [19–22]. In these nanocomposites, graphene acted as an efficient electron acceptor to enhance the charge transfer and thus reduces the electron-hole pair recombination in the composite to improve the photocatalytic activity. It has been also demonstrated that in TiO_2 -graphene based nanocomposites, the less defective graphene displayed a significantly larger enhancement in CO_2 photoreduction under visible light irradiation. This effect has been assigned to the superior mobility of charge carriers on graphene sheets, facilitating the diffusion of photoexcited electrons to reactive sites [19]. ZnO-rGO prepared through a simple hydrothermal process showed enhanced photocatalytic activity for the reduction of CO_2 to CH_3OH . The yield of CH_3OH on ZnO-rGO was found to be five times higher than that obtained on pure ZnO. The ZnO-rGO nanocomposites possessed excellent cycling property [23]. Similarly, loading copper nanoparticles (Cu NPs) of 4–5 nm in diameter onto GO was found to enhance the photocatalytic performance of pristine GO for the reduction of CO_2 to acetaldehyde and methanol under visible light irradiation [24]. An enhancement factor greater than 60 times has been achieved using Cu NPs/GO compared to pristine GO. The increased photocatalytic activity of GO was attributed to the suppression of electron-hole pair recombination, reduction of GO's band gap and modification of its work function. Independently, An et al. examined the catalytic activity of Cu_2O microparticles supported on reduced graphene oxide ($\text{Cu}_2\text{O}/\text{rGO}$) for the photoreduction of CO_2 [25]. The obtained results clearly demonstrated the positive effect of coupling Cu_2O with rGO, as witnessed by two times enhancement of the Cu_2O photoreduction activity and selectivity with CO being the only reduction product. Moreover, incorporation of Cu_2O onto rGO sheets improved the photocatalyst stability remarkably, holding a great potential for CO_2 conversion in a sustainable manner.

GO and rGO functionalization with visible light absorbing transition metal complexes and clusters was recently adopted as a very effective strategy to enhance the photocatalytic performance of GO (rGO) for the reduction of CO_2 into methanol. Various transition metal complexes such as ruthenium trinuclear polyazine complex [26], cobalt(II) phthalocyanine [27], copper(II) complex [28], heteroleptic ruthenium complex [29] and hexamolybdenum clusters [30] have been anchored to GO or nitrogen-doped graphene and successfully used as effective photocatalysts for the reduction of CO_2 into methanol under visible light irradiation.

In this work, we propose a multicomponent photocatalyst consisting of reduced graphene oxide wrapped core-shell structured magnetically separable $\text{rGO}@ \text{CuZnO}@\text{Fe}_3\text{O}_4$ microspheres and its successful application for the photoreduction of CO_2 into

methanol under visible light irradiation. The present photocatalyst takes advantage of the high photocatalytic efficiency, high electron mobility, low cost and environmental sustainment of ZnO, the high surface area and superior charge carrier mobility of rGO and the magnetic properties of the core Fe_3O_4 microspheres to design high efficient and recyclable photocatalyst for the selective reduction of CO_2 into methanol under visible light irradiation.

2. Experimental section

2.1. Materials

Iron (III) chloride hexahydrate (97%), polyethylene glycol (PEG, Mw~4000), sodium acetate trihydrate (99%), ethylene glycol (99.8%), zinc acetate dihydrate (98%), copper acetate monohydrate (99%), lithium hydroxide (LiOH, 98%), oleic acid (99%), graphite flakes, sodium nitrate (NaNO_3 , 99%), potassium permanganate (KMnO_4 , 99%), dimethylformamide (DMF) and 3-aminopropyltrimethoxysilane (APTMS, 99%) were purchased from Aldrich; they were of analytical grade and used without further purification. Sulfuric acid (H_2SO_4), ethanol, hydrochloric acid (HCl, 37%), and hydrogen peroxide (H_2O_2) were of analytical grade and purchased from Merck. All other chemicals were of A.R. grade and used without further purification. Graphene oxide (GO) was synthesized by a modified Hummer's method [31].

2.2. Techniques used for the chemical and structural characterizations

The rough structure of materials was determined by scanning electron microscopy using FE-SEM (Jeol Model JSM-6340F). The ultrafine structure of materials was determined with High Resolution Transmission Electron Microscopy (HR-TEM) using FEI-Tecnaig² Twin TEM operating at an acceleration voltage of 200 kV. It is equipped with a STEM module and an Oxford SDD 80 mm² EDX detector. The images were taken with a GATAN Orius 200D for medium resolution and a GATAN UltraScan 1000 for high resolution. The samples for TEM were prepared by depositing a very dilute aqueous suspension of material on carbon coated nickel grid. Fourier transform infrared (FTIR) spectra were recorded on a Perkin-Elmer spectrum RX-1 IR spectrophotometer using potassium bromide window. Raman spectra of materials were collected at room temperature using a Raman Microprobe (HR-800 Jobin-Yvon) with 532 nm Nd-YAG excitation source. The phase structure and crystalline nature of material were determined by powder X-ray diffraction pattern obtained on a Bruker D8 Advance diffractometer at 40 kV and 40 mA with Cu K_α radiation ($\lambda=0.15418$ nm). Surface properties of materials like Brunauer-Emmet-Teller surface area (S_{BET}), Barret-Joiner-Halenda (BJH) porosity, pore volume etc. were calculated by N_2 adsorption-desorption isotherm at 77 K by using VP. Micromeritics ASAP2010.

XPS measurements were obtained on a KRATOS-AXIS 165 instrument equipped with dual aluminum-magnesium anodes by using $\text{MgK}\alpha$ radiation ($h\nu=1253.6$ eV) operated at 5 kV and 15 mA with pass energy of 80 eV and an increment of 0.1 eV. To overcome the charging problem, a charge neutralizer of 2 eV was applied and the binding energy of C_{1s} core level ($\text{BE}\approx 284.6$ eV) of adventitious hydrocarbon was used as a standard. The XPS spectra were fitted by using a nonlinear square method with the convolution of Lorentzian and Gaussian functions, after a polynomial background was subtracted from the raw spectra. UV-vis absorption spectra of GO in water (0.03 mg/mL) was collected on Perkin Elmer lambda-19 UV-VIS-NIR spectrophotometer using a 10 mm quartz cell. While for all other solid samples UV-vis spectra were recorded

by making a 1 mm thick pellets using BaSO_4 as reference. The 333.6 nm line of a 30 mW argon laser was used to record the photoluminescence (PL) spectra at room temperature. The laser beam was focused with a 100 mm focal length lens to produce a 30- μm diameter spot on the sample. The incident power on the sample was 10 mW. The luminescence emitted by the specimen was analyzed by a triple monochromator Dilor RT 30, and then detected using a Hamamatsu R943 photomultiplier in the photon counting mode. Thermal stability and degradation pattern of material were determined by thermogravimetric analysis (TGA) using a thermal analyzer TA-SDT Q-600. Analysis was carried out in the temperature range of 40–800 °C under nitrogen flow with a heating rate of 10 °C/min. Elemental analysis of sample was done on Thermo Scientific™ FLASH 2000 Series CHNS/O Analyzer.

2.3. Synthesis of Fe_3O_4 magnetic microspheres **1**

Magnetic Fe_3O_4 microspheres were synthesized by following a literature method [32]. Briefly to 40 mL ethylene glycol were added, 1.35 g (5.0 mmol) of iron (III) chloride hexahydrate, 1.0 g PEG_{400} and 3.6 g (26.5 mmol) sodium acetate trihydrate and the resulting mixture was stirred to form a clear solution. The obtained solution was transferred in a Teflon lined autoclave and heated at 180 °C for 19 h. After cooling to room temperature, the obtained microspheres were collected by external magnet, washed with water and ethanol, and dried at 60 °C for 8 h.

2.4. Synthesis of core-shell $\text{CuZnO@Fe}_3\text{O}_4$ microspheres **2**

The core-shell $\text{CuZnO@Fe}_3\text{O}_4$ microspheres having 1 wt% Cu were synthesized by a modification of literature procedure [33]. In a typical synthesis, 0.1 g of Fe_3O_4 microspheres, 0.88 g (4.0 mmol) zinc acetate dihydrate and 0.027 g (0.135 mmol) of copper acetate were added to 80 mL ethanol and sonicated for 10 min. The obtained solution was heated at 70 °C with stirring for 1 h. A separate solution of LiOH was made by dissolving 0.23 g (10.0 mmol) of LiOH in 80 mL of EtOH with stirring for 20 min. Then the ethanolic LiOH solution was added drop-wise to Zn^{2+} , Cu^{2+} and Fe_3O_4 microspheres containing solution at 0 °C with vigorous stirring. The obtained suspension was sonicated for 5 min and 0.25 mL oleic acid as surfactant was added to prevent agglomeration. The resulting core-shell microspheres were collected by an external magnet and washed thoroughly with water and ethanol followed by drying at 50 °C under vacuum for 5 h for removing the excess of oleic acid from the surface of microspheres. For crystallization of the ZnO shell, the microspheres were subjected to calcination at 500 °C for 8 h. Calcination at higher temperature allowed the removal of residual oleic acid from the surface of microspheres. However, as an effect of higher temperature, the surface –OH group were also removed; thus, for restoration of –OH groups, microspheres were stirred in water for 6 h and dried at 70 °C for 12 h. Furthermore, for the comparison purpose, $\text{ZnO@Fe}_3\text{O}_4$ microspheres without Cu loading were synthesized and tested for photoreduction of CO_2 under identical conditions.

2.5. Amino-functionalization of $\text{CuZnO@Fe}_3\text{O}_4$ microspheres

The surface of $\text{CuZnO@Fe}_3\text{O}_4$ microspheres was functionalized with amino functionalities by refluxing 0.1 g $\text{CuZnO@Fe}_3\text{O}_4$ and 0.1 mL 3-aminopropyl trimethoxysilane (APTMS) in 25 mL ethanol for 24 h. The amino functionalized $\text{CuZnO@Fe}_3\text{O}_4$ microspheres were separated by an external magnet and washed with ethanol in order to remove unreacted APTMS.

2.6. Synthesis of $\text{GO@CuZnO@Fe}_3\text{O}_4$ microspheres **3**

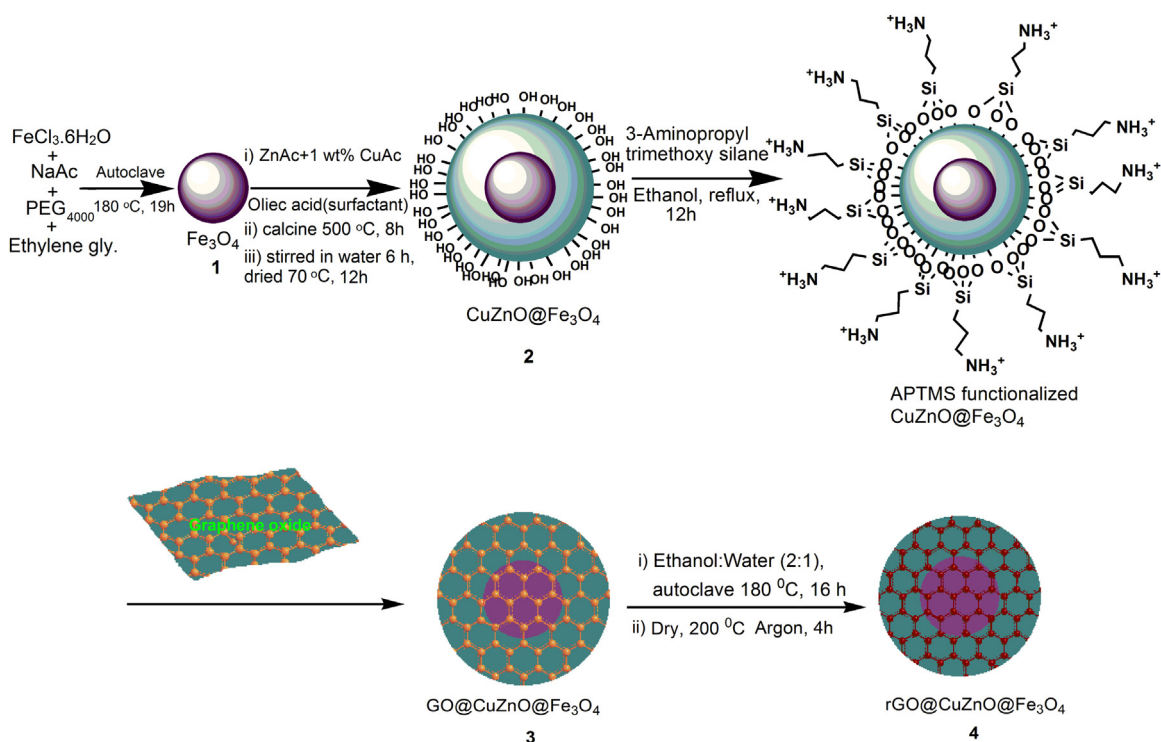
The GO was wrapped on the surface through ionic interaction between positively charged amino functionalized $\text{CuZnO@Fe}_3\text{O}_4$ **2** microspheres and negatively charged GO sheets. In a detail experimental procedure, APTMS functionalized $\text{CuZnO@Fe}_3\text{O}_4$ microspheres dispersed in 50 mL ethanol were added to 20 mL (0.2 mg/mL) GO aqueous suspension under vigorous stirring at pH 6. After stirring for 30 min, the GO wrapped $\text{CuZnO@Fe}_3\text{O}_4$ microspheres ($\text{GO@CuZnO@Fe}_3\text{O}_4$) **3** were separated by external magnet, washed with deionized water and dried at 40 °C. Similarly, for a control experiment the corresponding $\text{GO@ZnO@Fe}_3\text{O}_4$ microspheres without copper were synthesized.

2.7. Preparation of $\text{rGO@CuZnO@Fe}_3\text{O}_4$ **4**

The $\text{GO@CuZnO@Fe}_3\text{O}_4$ microspheres **4** were subjected to hydrothermal treatment for effective reduction of GO into reduced graphene oxide. In detail, $\text{GO@CuZnO@Fe}_3\text{O}_4$ microspheres were dispersed in 30 mL ethanol: water (2:1) solution and transferred in a Teflon-lined stainless steel autoclave. The mixture was heated at 180 °C for 16 h. After cooling to room temperature the suspension was transferred in a beaker and the particles were collected by a magnet, washed with ethanol and dried at 60 °C. Then the particles were calcined at 200 °C for 4 h under argon atmosphere to remove organic impurities. Similarly, rGO@CuZnO and $\text{rGO@ZnO@Fe}_3\text{O}_4$ microspheres were synthesized for comparing the photocatalytic performance.

2.8. Photocatalytic reduction of carbon dioxide

The photocatalytic activity of $\text{rGO@CuZnO@Fe}_3\text{O}_4$ **4** and other counterpart components was evaluated using visible light ($\lambda > 400 \text{ nm}$) of a 20 W white cold LED light (Model No. HP-FL-20W-F, Hope LED Opto-Electric Co., Ltd. Specification-Chip: KWE, Bridgelux, Epistar COB LED, 20W, Lum output: 80–90 lm/w, Beam angle: 120 degree and wavelength $\lambda > 400 \text{ nm}$ (maximum at 515 nm). In a 100 mL borosil cylindrical vessel (5 cm dia.), a mixture of DMF and water (45 mL/5 mL) was charged and purged with nitrogen to remove dissolved gases. After that CO_2 gas was purged in this mixture for 30 min to saturate the solution. Finally, the reaction vessel was charged with 100 mg of catalyst and sealed with a rubber septum. The reaction vessel was placed in a box equipped with magnetic stirrer to prevent from external interference and irradiated with visible light by using LED. The intensity of light on the vessel, measured by intensity meter, was equal to 85 W m^{-2} . To check the progress of reaction after every 2 h, a sample was withdrawn from the reaction vessel with the help of a needle, and the catalyst was separated by syringe filter (0.2 μm PTFE, 13 mm diameter). The photoreaction products were identified with GC-FID (Varian CP3800 by using CP Sil 24CB LOW BLEED/MS column, flow rate: 0.5 mL min^{-1} , injector temperature: 250 °C, FID detector temperature: 275 °C). Gaseous products were identified by withdrawing 20 μL sample and analyzed with GC–TCD and GC–FID (Agilent 7890A GC system) using capillary column (RGA, refinery gas analyzer) at the flow rate (H_2 : 35 mL min^{-1} , air: 350 mL min^{-1} , makeup flow: 27 mL min^{-1} , for TCD reference flow: 45 mL min^{-1} , Helium flow: 2 mL min^{-1}), injector temperature: 220 °C, TCD detector temperature and FID detector temperature: 220 °C. To quantify and check the linear response of GC with different concentrations, a calibration curve was plotted by injecting 1 μL of 50 ppm, 100 ppm, 200 ppm and 500 ppm methanol in GC–FID. Three blank reactions were carried out to confirm that the produced methanol was originated from the CO_2 photoreduction. For the first blank reaction all conditions were kept identical except without adding photocatalyst to the reaction vessel. In the second blank reaction,



Scheme 1. Synthesis of rGO@CuZnO@Fe₃O₄ microspheres **4**.

photocatalyst and other parameters were same, except the reaction was performed in the dark. While for the third blank reaction, N₂ was purged in place of CO₂, under otherwise identical conditions. In all three blank reactions no product was formed which confirmed that the produced methanol was the product of CO₂ photoreduction.

3. Results and discussion

3.1. Synthesis and characterization of the photocatalyst

The synthetic route of rGO@CuZnO@Fe₃O₄ microspheres **4** is depicted in Scheme 1. At first Fe₃O₄ microspheres **1** were synthesized by following the already known hydrothermal method using iron(III) chloride hexahydrate as precursor [32]. The core-shell structured CuZnO@Fe₃O₄ **2** microspheres were prepared by precipitation of zinc acetate and copper acetate solution on Fe₃O₄ microspheres **1** using lithium hydroxide and oleic acid as surfactant for maintaining spherical geometry of the microspheres, followed by calcination at 500 °C for 8 h [33]. The copper content was maintained to 1 wt% of ZnO. Calcination at higher temperature removed the remaining oleic acid as well as surface –OH groups. The surface –OH groups were restored and maintained in plenty by stirring the sample in water followed by drying at 70 °C. Thus obtained microspheres **2** were subsequently treated with APTMS to get amino functionalized **2**. In the next step, graphene oxide (GO) [31] was wrapped on the surface of –NH₂ functionalized CuZnO@Fe₃O₄ microspheres through electrostatic interactions [34]. Subsequently GO wrapped microspheres **3** were subjected to hydrothermal treatment at 180 °C for 16 h to get targeted photocatalyst rGO@CuZnO@Fe₃O₄ **4**. Thus obtained **4** was finally calcined at 200 °C for 4 h under argon atmosphere to remove the residual organic part.

The surface morphology and core-shell structure of microspheres were investigated by FE-SEM imaging (Fig. 1). All Fe₃O₄ particles **1** revealed a spherical morphology with a rough surface;

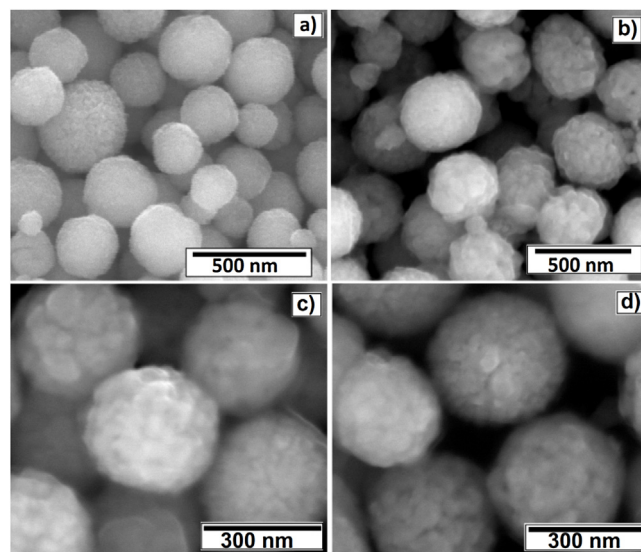


Fig. 1. FE-SEM images of a) Fe₃O₄ **1**, b) CuZnO@Fe₃O₄ **2**, c) GO@CuZnO@Fe₃O₄ **3** and d) rGO@CuZnO@Fe₃O₄ **4**.

their diameter varied between 200 and 250 nm (Fig. 1a) [32–34]. After coating with CuZnO the particles remained spherical with some rough outgrowths on the surface and their mean diameter increased up to 300 nm (Fig. 1b). Such outgrowths were probably generated during ZnO crystallization upon calcination at 500 °C for 8 h. The SEM image of GO@CuZnO@Fe₃O₄ **3** revealed similar morphology like CuZnO@Fe₃O₄ along with wrapped GO sheets on the surface (Fig. 1c). The structure of rGO@CuZnO@Fe₃O₄ **4** remained intact after GO reduction to rGO (Fig. 1d).

EDX analysis showed that the samples contained only Fe and O elements for Fe₃O₄ **1**; Zn, Fe and O elements for CuZnO@Fe₃O₄ **2**, Cu was not detected may be due to its low concentration in the

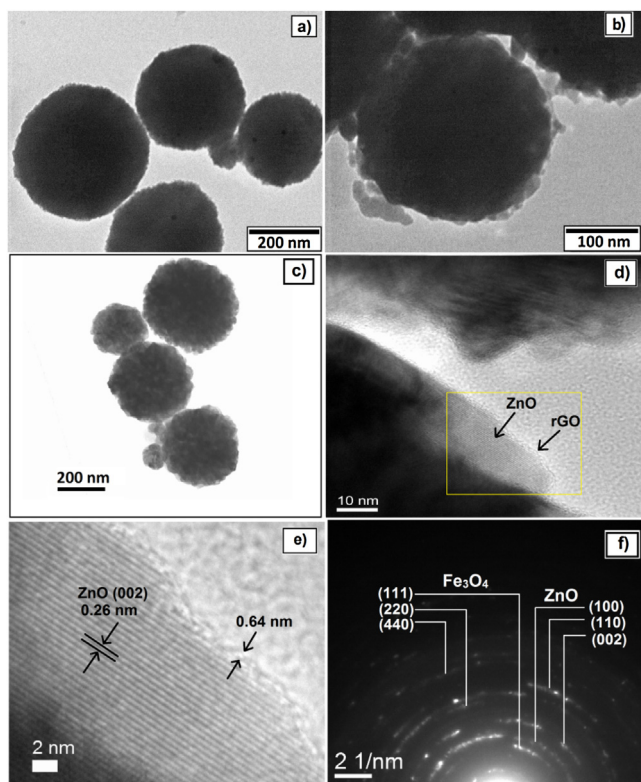


Fig. 2. Low resolution TEM images of a) Fe_3O_4 **1**, b) $\text{CuZnO@Fe}_3\text{O}_4$ **2**, c) $\text{rGO@CuZnO@Fe}_3\text{O}_4$ **4**, d) high magnification TEM of **4**, e) high resolution TEM and f) SAED pattern of $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres **4** showing diffraction patterns of Fe_3O_4 and ZnO.

material; C, Zn, Fe and O elements for samples **3** and **4** (See Fig. S1, Supplementary Information).

Fine structures of Fe_3O_4 **1**, $\text{CuZnO@Fe}_3\text{O}_4$ **2** and $\text{rGO@CuZnO@Fe}_3\text{O}_4$ **4** microspheres were examined using TEM imaging (Fig. 2). Low resolution TEM image of **1** exhibited spherical particles of 200–250 nm mean diameter (Fig. 2a). After coating with CuZnO, the surface of $\text{CuZnO@Fe}_3\text{O}_4$ **2** displayed a rough structure due to ZnO deposition, and the particle size slightly increased up to 300 nm (Fig. 2b). High resolution TEM image of $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres **4** showed that particles retained similar rough surface appearance as $\text{CuZnO@Fe}_3\text{O}_4$ **2** and their diameter were in the 200–300 nm range (Fig. 2c). The TEM

image of $\text{rGO@CuZnO@Fe}_3\text{O}_4$ **4** at higher magnification clearly evidenced the rough surface and lattice fringes of ZnO with the outer amorphous zone attributed to rGO (Fig. 2d) [35,36]. The TEM image of the **4** in the high resolution mode clearly exhibited the 1 D lattice plane with 0.26 nm interplanar distance (d-spacing) correlated to (002) plane of ZnO (Fig. 2e). In addition, a 0.64 nm of thickness amorphous zone correlated to rGO was visible (Fig. 2e). Selected area electron diffraction pattern of microspheres **4** gave diffraction spots corresponding to (111), (220) and (440) planes of Fe_3O_4 (JCPDS No. 19–0629) and (100), (110), and (002) planes of ZnO (JCPDS No. 36–1451) (Fig. 2f).

Scanning transmission electron microscopy (STEM) elemental mapping of $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres **4** is depicted in Fig. 3. It could be seen that O, Cu, Zn, Fe and C elements were homogeneously distributed in the material. The presence of an additional carbon zone was most likely due to carbon from the TEM grid. However, a higher concentration of carbon at the microsphere location confirmed the presence of rGO sheets. The presence of only C, Cu, Zn, Fe and O in the microspheres was furthermore corroborated by EDX analysis (Fig. S2). A high intensity of carbon was observed as a result of the carbon coated TEM grid.

Fig. 4 exhibits Raman spectra of $\text{GO@CuZnO@Fe}_3\text{O}_4$ **3**, $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres **4** along with GO and rGO spectra. The Raman spectrum of GO revealed specific D (1350 cm^{-1}) and G (1593 cm^{-1}) bands originated from out-plane vibration of sp^3 carbons presented in the form of defects and in plane vibration of sp^2 carbons in aromatic conjugated system of GO, respectively. The intensity ratio of D to G bands (I_D/I_G) was about 1.01, indicating the presence of many defects in GO in the form of epoxide and other oxygen carrying groups. For rGO the positions of D and G bands were found to be at 1349 and 1593 cm^{-1} , respectively and the integral intensity ratio of D to G bands (I_D/I_G) was equal to 0.41. The decrease of D band intensity was attributed to the removal of oxygen carrying functionalities during reduction step and restoration of aromatic conjugated system. For $\text{GO@CuZnO@Fe}_3\text{O}_4$, the D and G bands of GO were observed at 1351 and 1594 cm^{-1} , respectively with a more intense D band ($I_D/I_G = 1.13$), indicating a large number of defects specific of GO. The decrease of D band (1356 cm^{-1}) intensity in comparison to G band (1496 cm^{-1}) with $I_D/I_G = 0.87$ after hydrothermal treatment of $\text{GO@CuZnO@Fe}_3\text{O}_4$ **3** confirmed the successful reduction of GO to rGO in **4**.

In addition, the Raman spectrum of $\text{GO@CuZnO@Fe}_3\text{O}_4$ **3** displayed a peak at 664 cm^{-1} due to A_{1g} vibration mode of Fe_3O_4 [37] and a band at 325 cm^{-1} due to A_1 vibration mode of wurtzite ZnO [38]. Some peaks of Fe_3O_4 and ZnO overlapped. The same peaks

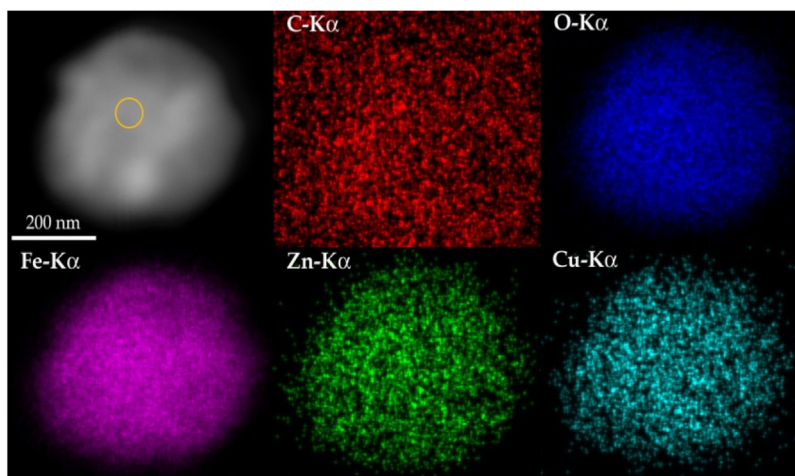


Fig. 3. STEM elemental mapping of $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres **4**.

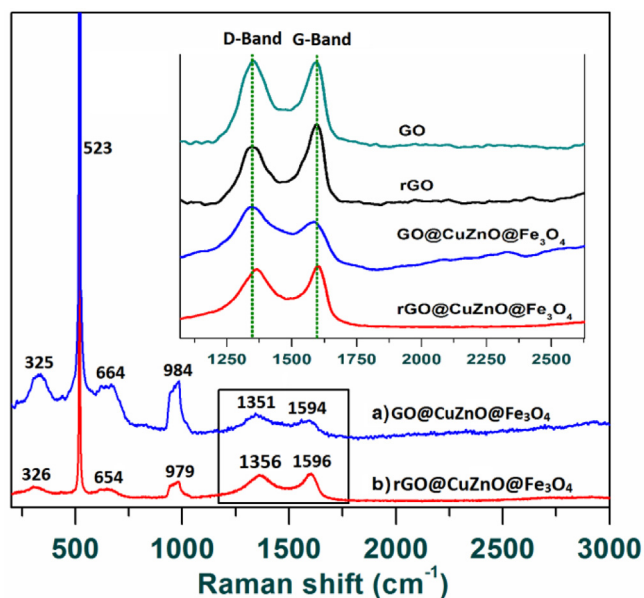


Fig. 4. Raman spectra of a) GO@CuZnO@Fe₃O₄ **3**, b) rGO@CuZnO@Fe₃O₄ **4**. The insets correspond to the D and G bands of GO and rGO. The sharp peak at 523 cm⁻¹ is due to silicon substrate.

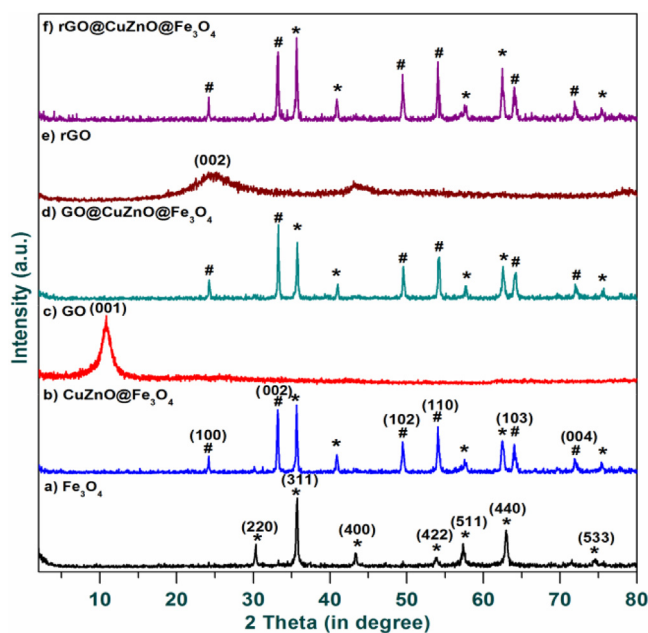


Fig. 5. XRD diffraction patterns of a) Fe₃O₄ (**1**), b) CuZnO@Fe₃O₄ (**2**), c) GO, d) GO@CuZnO@Fe₃O₄ (**3**), e) rGO and f) rGO@CuZnO@Fe₃O₄ (**4**).

were observed in the Raman spectrum of rGO@CuZnO@Fe₃O₄ **4**, indicating the presence of ZnO wurtzite and Fe₃O₄ magnetite in both samples (Fig. 4).

Powder X-ray diffraction patterns of Fe₃O₄ **1**, CuZnO@Fe₃O₄ **2**, GO, GO@CuZnO@Fe₃O₄ **3**, rGO and rGO@CuZnO@Fe₃O₄ **4** are depicted in Fig. 5. The XRD pattern of **1** showed characteristic Bragg peaks at 2θ values of 30.32° (220), 35.72° (311), 43.4° (400), 53.92° (422), 57.32° (511), 62.92° (440) and 74.56° (533) matching perfectly with face centered cubic inverse spinel structure of magnetite according to JCPDS No. 19-0629 (Fig. 5a). After coating with 1 wt% copper containing ZnO on Fe₃O₄ microspheres **1**, the core-shell structured microspheres **2** showed characteristic diffraction peaks of hexagonal wurtzite ZnO along with Fe₃O₄ diffraction peaks at 2θ : 24.20° (100), 33.16° (002), 49.48° (102), 54.04° (110), 64.12°

(103) and 71.84° (004) which was in well conformity with JCPDS No. 36-1451 (Fig. 5b).

No diffraction peak of copper was observed due to its low concentration in comparison to bulk material. Pristine GO gave a sharp peak at $2\theta = 11.3^\circ$ due to diffraction of (001) plane (Fig. 5c). However, no peak of GO was seen in the XRD diffractogram of **3** which indicated the presence of few GO layers in **3** (Fig. 5d). rGO gave a broad hump at $2\theta = 24.36^\circ$ due (002) carbon plane reflection (Fig. 5e), but this peak was absent in the XRD pattern of rGO@CuZnO@Fe₃O₄ **4** (Fig. 5f).

X-ray photoelectron spectroscopy (XPS) was used to follow the changes in the chemical composition at each stage of material modification (Fig. 6). Survey scans of microspheres **3** and **4** exhibited peaks at 285, 530, 400, 710, 1022 and 103 eV due to C_{1s}, O_{1s}, N_{1s}, Fe_{2p}, Zn_{2p} and Si_{2p}, respectively. The peak of copper due to its lower concentration may not be appearing in the XPS; however, its presence was confirmed by EDX (Fig. 3). The atomic percentages of different components are given in Table S1; the carbon atomic percentage was estimated to be 56.16% and 48.50% for microspheres **3** and **4**, respectively. Higher value of carbon concentration suggested the presence of carbon sheets on the surface of **4**. Furthermore, the presence of small amount of Si_{2p} and N_{1s} was detected due to the APTMS linker. C_{1s} high resolution XPS spectrum of microspheres **3** exhibited characteristic peak components of GO at 285.1, 286.6 and 289.0 eV due to C–C, C–O and –COOH groups, respectively with an additional peak at 283.7 eV due C–Si group (Fig. 6a). After hydrothermal reduction, the C_{1s} wide scan of microspheres **4** showed a decrease of –COOH peak intensity at 289.0 eV, suggesting the successful reduction of GO to rGO (Fig. 6b).

Fe_{2p} high resolution XPS spectrum of microspheres **4** displayed two characteristic peaks due to Fe_{2p_{3/2}} and Fe_{2p_{1/2}} at 711.5 and 724.8 eV, respectively, in concordance with literature values for Fe₃O₄ (Fig. 6c). Further, a satellite peak was observed at 718 eV attributed due to presence very thin oxide's layer of Fe₂O₃ on the surface of microspheres. In addition, the peaks at 1022.0 and 1043.1 eV due to Zn_{2p_{3/2}} and Zn_{2p_{1/2}}, respectively confirmed the presence of ZnO on the surface of microspheres **4** (Fig. 6d). We did not observe any peak due to Cu, because of its low concentration in comparison to ZnO in the composite.

UV–vis absorption spectra of solid samples are shown in (Fig. 7a). UV–vis absorption spectrum of magnetic Fe₃O₄ core and Fe₂O₃ present on surface of microspheres. Nanoparticles gave a broad absorption profile from 200 to 650 nm, due to d-orbital transitions of Fe₃O₄ [39–41]. ZnO synthesized by similar method gave a sharp absorption band at 370 nm and thus absorbs only in UV region. The core-shell CuZnO@Fe₃O₄ microspheres **2** displayed a broad absorption band from 200 to 560 nm similar to pure Fe₃O₄ but the intensity started to decrease after 560 nm due to the coating of ZnO. After wrapping of GO sheets on CuZnO@Fe₃O₄ microsphere **2**, the visible light absorption pattern slightly decreased in GO@CuZnO@Fe₃O₄ **3** as GO sheets could not absorb in visible region. Further for rGO wrapped rGO@CuZnO@Fe₃O₄ microspheres **4** the absorption pattern increased in visible region due to the synergistic effect.

The optical band gap of the Fe₃O₄ **1**, ZnO, CuZnO@Fe₃O₄ **2**, GO@CuZnO@Fe₃O₄ **3** and rGO@CuZnO@Fe₃O₄ **4** was calculated by extrapolating the rising segment of the UV spectrum to the abscissa at zero absorption with the help of the Tauc plot (Fig. 7b). Although Fe₃O₄ is semi-metal and doesn't possess a band gap at 300 K but during synthesis and oxidation in air a very thin layer of oxide (Fe₂O₃) was obtained on the surface of Fe₃O₄ microspheres. So, due to presence of Fe₂O₃ on the surface of Fe₃O₄ microspheres can work as semiconductor [42]. In our case a band gap value of **1** was found to be 1.16 eV which was low enough for visible light mediated transition. The value of the band gap of pure sample of ZnO was 3.20 eV (corresponding to 386 nm), which indicates that

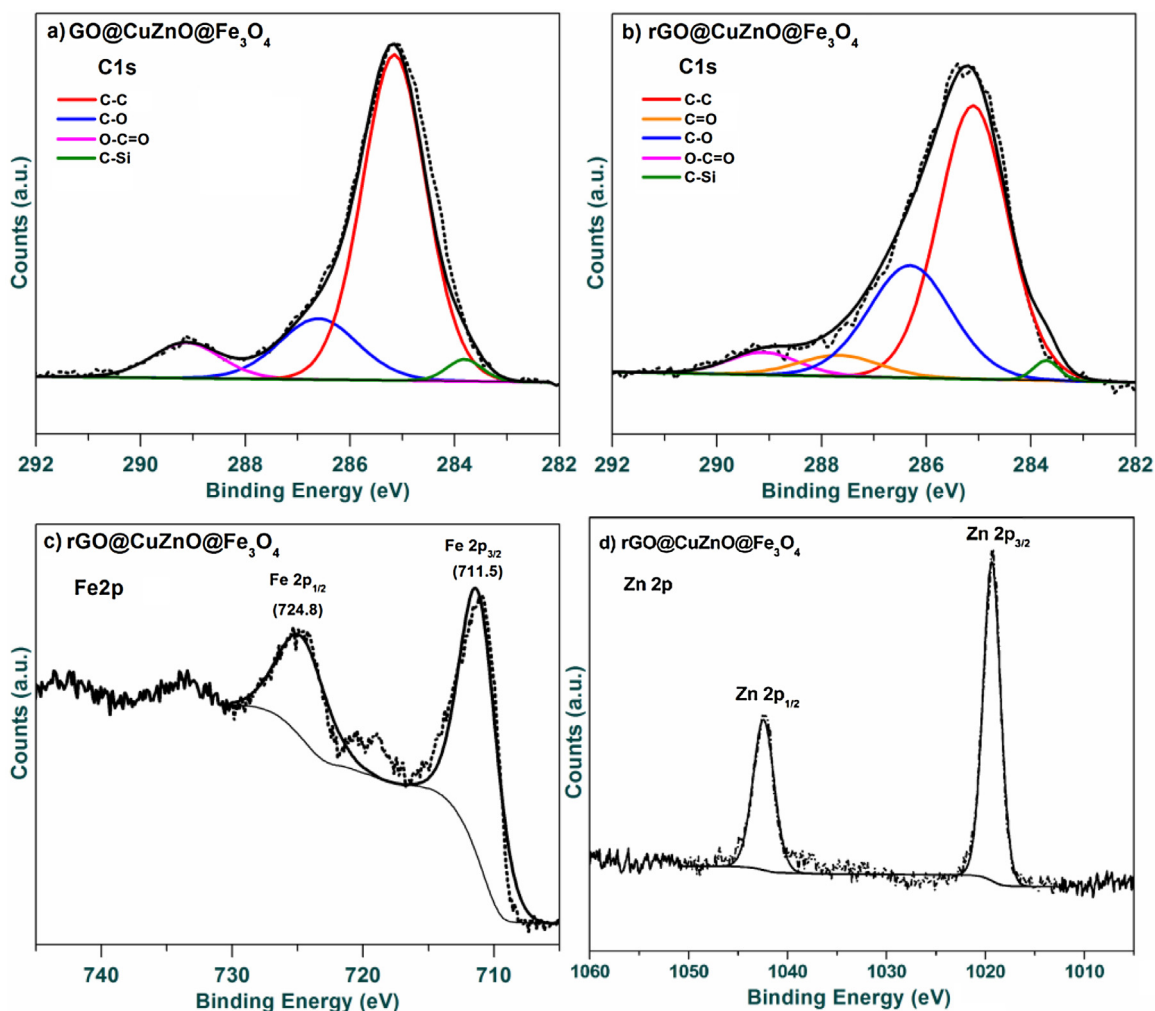


Fig. 6. C_{1s} high resolution XPS spectra of a) $GO@CuZnO@Fe_3O_4$ **3**, b) $rGO@CuZnO@Fe_3O_4$ **4**, c) Fe_{2p} and d) Zn_{2p} of $rGO@CuZnO@Fe_3O_4$ **4**.

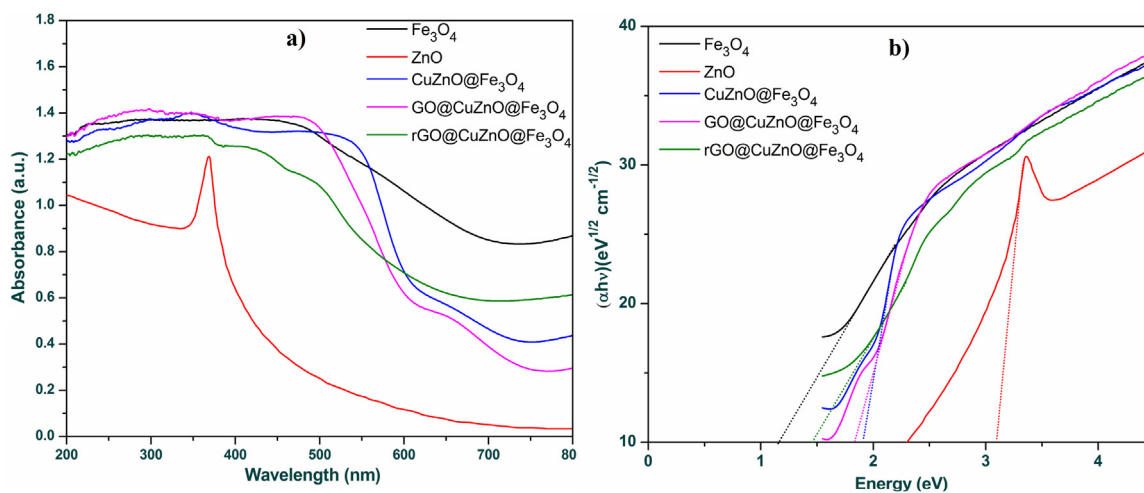


Fig. 7. UV-vis absorption spectra a) and Tauc plot showing band gap b) of Fe_2O_3/Fe_3O_4 **1**, $CuZnO@Fe_3O_4$ **2**, $GO@CuZnO@Fe_3O_4$ **3**, $rGO@CuZnO@Fe_3O_4$ **4**.

it can absorb only in the UV region. The obtained band gap value was found to be slightly lower than the reported value (3.3 eV) which is most likely due to the presence of some defects [43]. After the coating of ZnO on the Fe_3O_4 microspheres the value of the band gap increased to 1.9 eV associated with 651 nm wavelength. The GO wrapped particles $GO@CuZnO@Fe_3O_4$ **3** displayed a band

gap of 1.85 eV (corresponding wavelength: 669 nm). Hydrothermal treatment of **3** transformed wrapped GO to rGO and the band gap value decreased to 1.44 eV for $rGO@CuZnO@Fe_3O_4$ **4** which was correlated to 859 nm wavelength. The reduced band gap value for microspheres **4** confirmed their visible light absorption potential.

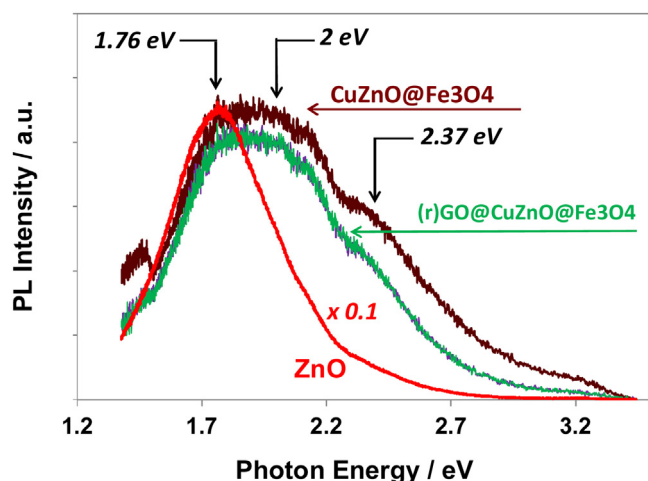


Fig. 8. Photoluminescence spectra of ZnO (red), CuZnO@Fe₃O₄ (dark red) **2**, GO@CuZnO@Fe₃O₄ **3** (green) and rGO@CuZnO@Fe₃O₄ **4** (blue). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

The PL spectra of the different samples are depicted in Fig. 8. Three important observations can be made: (i) the integral luminescence intensity ($I_{PL_{tot}}$) of ZnO is about 6 times that of the other three samples (CuZnO@Fe₃O₄, GO@CuZnO@Fe₃O₄ and rGO@CuZnO@Fe₃O₄), (ii) $I_{PL_{tot}}$ of CuZnO@Fe₃O₄ is about 1.3 times that of GO@CuZnO@Fe₃O₄ and of rGO@CuZnO@Fe₃O₄ which are identical, (iii) the spectral shape of ZnO is very different from the three other ones which are nearly identical. Its full width at half maximum (FWHM) is close to 0.5 eV, whereas it is as large as 0.8 eV in both GO@CuZnO@Fe₃O₄ **3** and rGO@CuZnO@Fe₃O₄ **4** and equal to 1 eV in CuZnO@Fe₃O₄. Furthermore, its shape is very asymmetric, with a main red PL band at 1.76 eV (704 nm). On the opposite, the shape of the other three PL spectra is more symmetrical. In fact, additionally to the red band at 1.8 eV, there is a red/orange band of similar intensity which is close to 2 eV (620 nm). Furthermore, a green band close to 2.4 eV is also visible. The spectra extend until about 3.4 eV. The green band close to 2.4 eV which appears as a hump in the PL spectra is certainly due to copper [44–46]. The origin of the red emission is more controversial since it was associated with zinc interstitials [47], zinc vacancies [48] or excess oxygen [49]. The red/orange emission is usually attributed to oxygen interstitial defects [50] or to excess oxygen [51].

3.2. Photocatalytic activity

The photocatalytic performance of rGO@CuZnO@Fe₃O₄ microspheres **4** and other components such as GO, rGO, ZnO, CuZnO, Fe₃O₄ **1**, ZnO@Fe₃O₄, CuZnO@Fe₃O₄ microspheres **2** and GO@ZnO@Fe₃O₄ **3** for CO₂ photoreduction was evaluated in DMF/water (45/5 mL) using a 20 W white cold LED as visible light source (see more details in experimental section). The results of these experiments are summarized in Table 1. For the identification and quantification of products, 1 μ L liquid sample was withdrawn at every 2 h interval and analyzed by GC–FID. The GC analysis showed that methanol was formed as the major liquid reaction product without any evidence for the formation of other possible reaction products like formic acid, ethanol, etc (Fig. S6). A calibration curve was plotted by injecting calculated amount of methanol in GC for quantification and checking linear response of GC toward various concentrations of methanol (Fig. S7). Gaseous products were analyzed by injecting 20 μ L of sample in GC–TCD and GC–FID by using RGA column. Gaseous product analysis showed that hydrogen was evolved in very small amount; other products if

any like methane, CO, etc were not detected under our experimental conditions. As methanol was the only liquid reaction product, a graph between methanol yield (μ mol g^{−1} cat) and reaction time was plotted to determine the catalyst performance as compared to other catalyst components (Figs. 10 and 11). After 24 h visible light irradiation of a CO₂ saturated DMF/water solution, the yield of methanol by using Fe₃O₄ microspheres **1**, CuZnO@Fe₃O₄ **2**, GO@CuZnO@Fe₃O₄ **3** and rGO@CuZnO@Fe₃O₄ **4** was determined to be 278, 858, 1749 and 2656 μ mol g^{−1} cat, respectively. The methanol formation rate (R_{MeOH}) for **1**, **2**, **3** and **4** was found to be 11.6, 35.7, 72.9 and 110.6 μ mol h^{−1} g^{−1} cat, respectively. The associated quantum yield (ϕ_{MeOH}) for methanol was estimated to be 0.0026, 0.0082, 0.0167 and 0.0253 mol einstein^{−1}, respectively.

The quantum yield of product was determined by using following formula:

$$\phi_{product} = [P] \times n / [\gamma]$$

Where, $\phi_{product}$ is quantum yield, [P] is moles of desired product, n is number of electrons required for reduction and $[\gamma]$ is moles of incident photons.

For methanol and hydrogen formation, the number of electrons required is 6 and 2, respectively.

The analysis of gaseous products provided the hydrogen yield as 16.2, 28.5 and 45.5 μ mol g^{−1} cat for CuZnO@Fe₃O₄ **2**, GO@CuZnO@Fe₃O₄ **3** and rGO@CuZnO@Fe₃O₄ **4**, respectively with formation rate (R_{H_2}) of 0.6, 1.2 and 1.9 μ mol h^{−1} g^{−1} cat, respectively.

The catalytic selectivity was determined by:

$$CS\% = [P] \times 100 / [P_{total}]$$

Where CS% is the catalytic selectivity in%, [P] is moles of desired product; $[P_{total}]$ is moles of total reaction products.

The catalytic selectivity in percentage (CS%) of Fe₃O₄ microspheres **1**, CuZnO@Fe₃O₄ microspheres **2**, GO@CuZnO@Fe₃O₄ microspheres **3** and rGO@CuZnO@Fe₃O₄ microspheres **4** for methanol was determined to be 100, 94.14, 98.39 and 98.31, respectively.

The efficiency of the developed rGO@CuZnO@Fe₃O₄ photocatalyst **4** for CO₂ photoreduction was much higher than those reported in the literature for different photocatalysts. A comparison of the photocatalytic performance of **4** with the state of the art photocatalysts for CO₂ reduction is presented in Table S2.

As can be seen in Table 1 only a very small amount of methanol can be detected for rGO due to the absence of band gap (zero band gap) [52] so it was unable to generate electron and hole pairs which are essential for photoreduction of CO₂. However, the origin of small amount of methanol (117 μ mol g^{−1} cat) can be explained on the basis of residual oxygen functionalities which remained after the reduction of GO to rGO. Owing to the presence of sp² and sp³ carbons on its surface, rGO acted like a semiconductor. The sp² hybridized carbon containing domains due to the free mobility of electrons behave like a conduction band while sp³ carbon containing domains due to tightly held electrons behave like a valence band. But the number of such sp² and sp³ domains in rGO is less in comparison to GO therefore poor yield of methanol was obtained. The band gap value of GO was found to be in the range of 2.8–3.6 eV due to the uneven oxidation which is in well concordance with literature values [53]. ZnO has larger band gap (3.3 eV) which is high enough to prevent any visible light mediated transition [43]. However, a small amount of methanol (211 μ mol g^{−1} cat) may be originated due to the presence of some defects in ZnO. Addition of copper enhances the photocatalytic performance of ZnO by capturing electrons from the conduction band of ZnO which delayed the charge recombination process, so the electron availability for the reduction of CO₂ was increased [54,55]. To evaluate the optimal

Table 1
Photocatalytic reduction of CO₂ by using different catalysts.^a

Entry	Catalyst	Reactant	Methanol Yield ($\mu\text{mol g}^{-1} \text{cat}$)	Quantum Yield (mole einstein ⁻¹)
1	Blank	CO ₂	Nil ^b	Nil
2	Fe ₃ O ₄ 1	CO ₂	278	0.0026
3	ZnO	CO ₂	211	0.0020
4	1 wt% CuZnO	CO ₂	383	0.0036
5	GO	CO ₂	470	0.0044
6	rGO	CO ₂	117	0.0011
7	rGO@CuZnO	CO ₂	648	0.0061
8	ZnO@Fe ₃ O ₄	CO ₂	524	0.0049
9	CuZnO@Fe ₃ O ₄ 2	CO ₂	858	0.0082
10	GO@ZnO@Fe ₃ O ₄	CO ₂	942	0.0089
11	rGO@ZnO@Fe ₃ O ₄	CO ₂	1124	0.0106
12	GO@CuZnO@Fe ₃ O ₄ 3	CO ₂	1749	0.0167
13	rGO@CuZnO@Fe ₃ O ₄ 4	CO ₂	2656	0.0253
		N ₂	Nil	Nil
		CO ₂	Nil	Nil

^a Reaction conditions: catalyst: 100 mg, solvent: DMF/water (45/5), visible light irradiation using 20 W LED ($\lambda > 400 \text{ nm}$), time: 24 h.

^b Without catalyst.

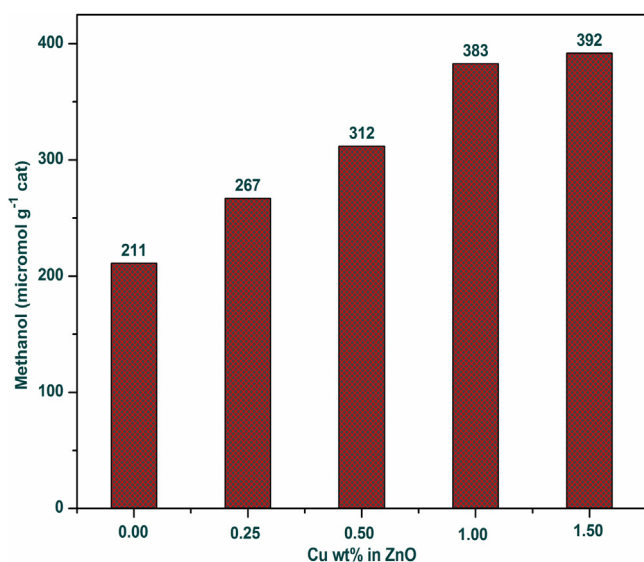


Fig. 9. Methanol yield using 0.25, 0.50, 1.00 and 1.50 wt% of copper loaded ZnO (CuZnO).

loading of Cu we synthesized a number of CuZnO composites having 0.25, 0.5, 1.0 and 1.5 wt% of copper. It can be seen from Fig. 9 that the yield of methanol increased with increasing the copper loading from 0.25 to 1.0 wt%, however further increase of Cu content did not influence the photocatalytic performance. The yield of methanol by using 0.25, 0.50, 1.0 and 1.5 wt% of copper loaded ZnO (CuZnO) was found to be 267, 312, 383 and 392 $\mu\text{mol g}^{-1} \text{cat}$, respectively. The enhancement in the methanol yield with copper content was assumed to be due to the trapping of photogenerated electrons which resulted in reduced electron-hole pair recombination. Based on these results we have chosen 1.0 wt% CuZnO as the optimal for coating on Fe₃O₄ microspheres **1**.

In case of Fe₃O₄ microspheres **1**, a very poor yield of methanol 278 $\mu\text{mol g}^{-1} \text{cat}$ was obtained due to presence of Fe₂O₃ layer on microspheres (Fig. 10b). In order to check the synergistic effect of the components in the designed composite **4** on CO₂ photoreduction, we synthesized all the possible combinations such as ZnO@Fe₃O₄, GO@ZnO@Fe₃O₄ and rGO@ZnO@Fe₃O₄ and evaluated their performance for the photoreduction of CO₂ under described experimental conditions. The yield of methanol by using ZnO@Fe₃O₄, GO@ZnO@Fe₃O₄ and rGO@ZnO@Fe₃O₄ microspheres as photocatalysts was found to be 524, 942 and 1124 $\mu\text{mol g}^{-1} \text{cat}$, respectively after 24 h of visible irradiation. For ZnO@Fe₃O₄ micro-

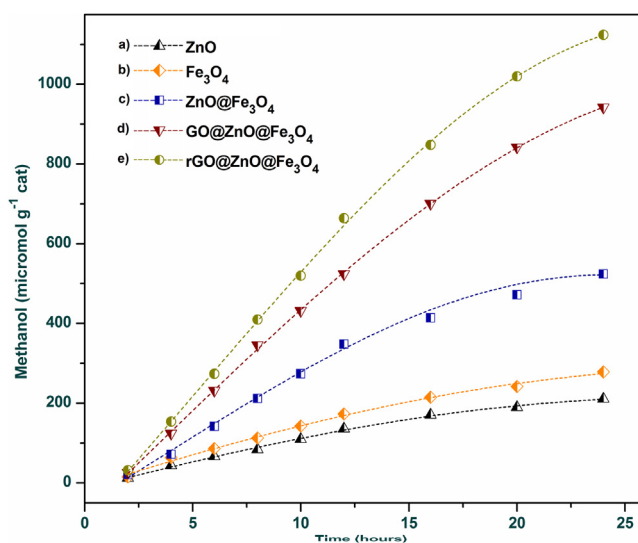


Fig. 10. CO₂ conversion to methanol for a) ZnO, b) Fe₃O₄ **1** c) ZnO@Fe₃O₄ microspheres, d) GO@ZnO@Fe₃O₄ microspheres, and e) rGO@ZnO@Fe₃O₄ microspheres.

spheres the enhancement of yield can be explained by the efficient transfer of photogenerated electrons and holes of Fe₂O₃/Fe₃O₄ to the conduction and valence band of ZnO, respectively (Fig. 10c). Further, the wrapping of GO on ZnO@Fe₃O₄ microspheres enhances the yield of methanol which is believed to be due to the transfer of electrons from CB of ZnO to GO sheets (Fig. 10d). Reduction of wrapped GO to rGO further enhances the yield of methanol because of the restoration of sp² hybridized aromatic system in rGO which provides better electron mobility as well as charge separation in comparison to GO where the presence of sp³ carbons restricts the free movement of the electrons on the GO sheets (Fig. 10e). Furthermore, the yield of methanol by using rGO@CuZnO as photocatalyst was found to be 648 $\mu\text{mol g}^{-1} \text{cat}$ which confirms that the coating of rGO played an important role in facilitating the charge separation.

As shown in the previous section that the presence of copper enhanced the photocatalytic performance and methanol yield. To establish the promoting effect of copper, we compared the photocatalytic activity of **2** and **3** with **4** (Fig. 11). No methanol was formed in the blank reaction in the absence of photocatalyst (Fig. 11a). However, the methanol yield using 1 wt% CuZnO was found to be 383 $\mu\text{mol g}^{-1} \text{cat}$ (Fig. 11b). The methanol yield for CuZnO@Fe₃O₄ microspheres **2** was determined to be 858 $\mu\text{mol g}^{-1} \text{cat}$ (Fig. 11c). The increase in yield was attributed to the transfer of electrons and holes from Fe₂O₃/Fe₃O₄ to the CB and VB of ZnO followed

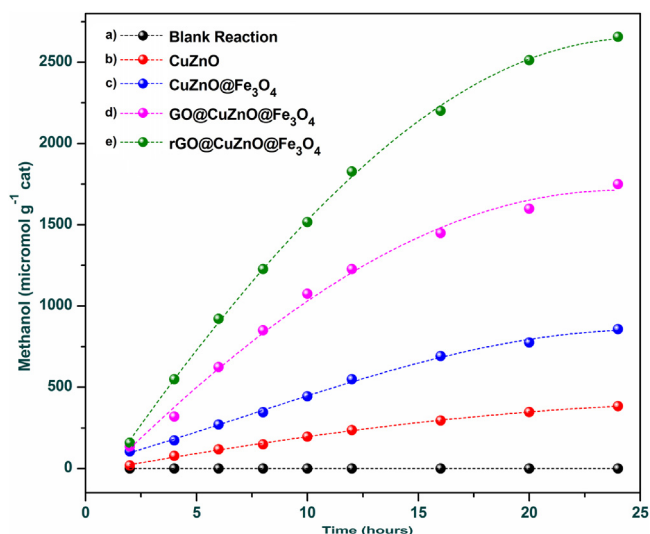


Fig. 11. Yield of methanol, a) blank reaction, b) using 1 wt% CuZnO , c) $\text{CuZnO@Fe}_3\text{O}_4$ 2, d) $\text{GO@CuZnO@Fe}_3\text{O}_4$ 3, and e) $\text{rGO@CuZnO@Fe}_3\text{O}_4$ 4.

by electron capturing by Cu, which prevents charge recombination. Because, CuZnO was coated on Fe_3O_4 microspheres by *in situ* method, we assume that copper was evenly distributed in the ZnO layer of the composite. XPS analysis of the nanocomposite did not show any peak of the Cu due to its low concentration in comparison to ZnO, which confirms that Cu is not limited to ZnO surface but it is also present in the bulk volume of ZnO (Fig. 6 and Table S1). Moreover, STEM elemental mapping and EDX analysis of microspheres 4 clearly showed the presence of Cu because of the deeper penetration by electron beam, so more Cu atoms are encountered in path, confirming that Cu was homogeneously distributed in the ZnO layer (Figs. 3 and S2). Despite that the Cu embedded in ZnO also takes part in electron capturing process, it does not participate in electron transfer to the GO/rGO due to deactivation of charge by the process of volume recombination. Only the Cu present on the ZnO surface can transfer captured electrons to rGO. The GO wrapped $\text{GO@CuZnO@Fe}_3\text{O}_4$ microspheres 3 afforded $1749 \mu\text{mol g}^{-1} \text{cat}^{-1}$ of methanol which was much higher than 2 due to the transfer of electrons to GO surface (Fig. 11d). The highest yield of methanol was obtained for $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres 4 up to $2656 \mu\text{mol g}^{-1} \text{cat}^{-1}$ because of better mobility of electrons on reduced graphene oxide surface which maintains the sufficient distance between the electrons and thus slows down charge recombination process (Fig. 11e).

The synthesized $\text{rGO@CuZnO@Fe}_3\text{O}_4$ (4) could be easily recovered by using external magnet for recycling experiments. Fig. 12a and b showed that the catalyst was well dispersed in absence of any magnetic field, while after exposure to 1 T magnet for 1 min the coloured solution became nearly transparent.

Furthermore, we performed recycling experiments in order to check the reusability of the $\text{rGO@CuZnO@Fe}_3\text{O}_4$ 4 photocatalyst. After completion of the reaction, the photocatalyst was separated by using a magnet, washed with water and reused for subsequent CO_2 photoreduction experiment. It can be seen from Fig. 12 that there was no significant loss in activity even after five recycling experiments. However, a slight decrease in the photocatalytic performance after few recycling runs was attributed to removal of some rGO sheets from the surface of 4 under vigorous stirring condition. To confirm the loss of rGO sheets from 4, we performed elemental analysis of freshly synthesized and recovered photocatalyst after fifth run to determine the carbon content. The carbon content of freshly synthesized 4 was 3.67 wt% while after five recycling the carbon content was 3.43 wt%. This small decrease

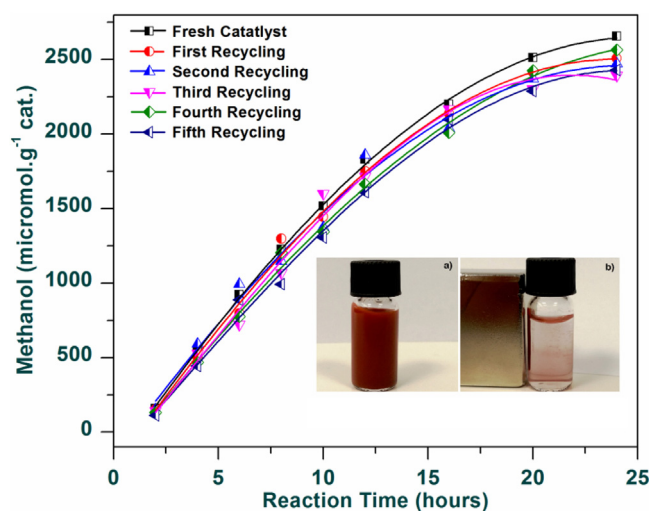
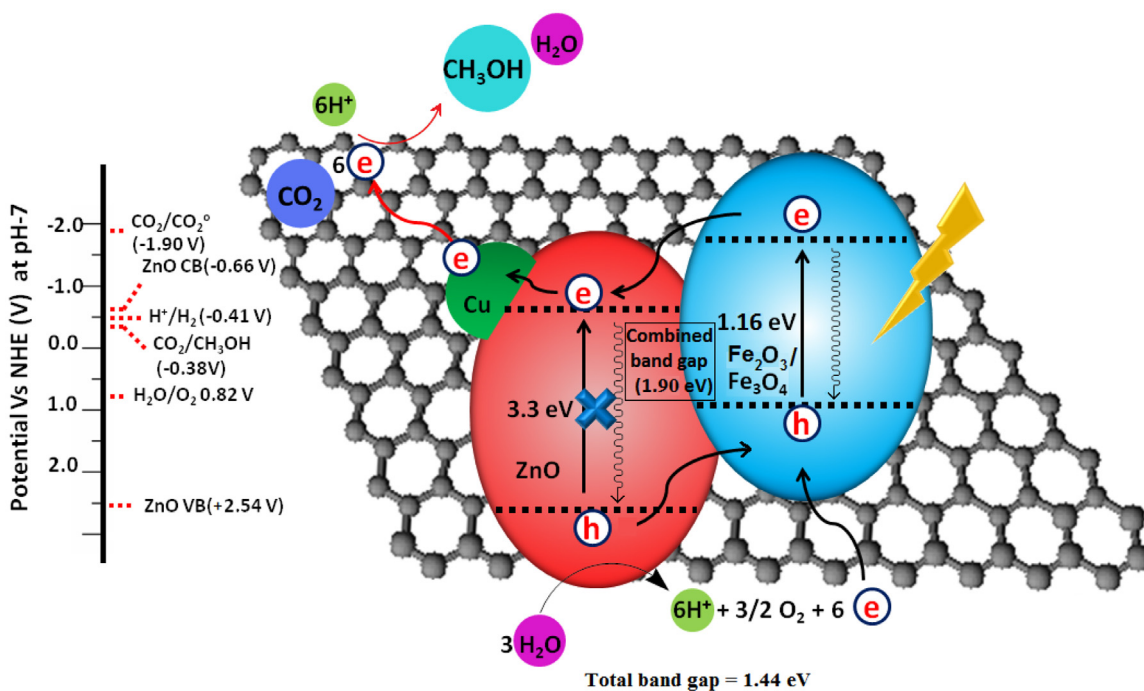


Fig. 12. Recycling experiments using $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres 4. Inset showing magnetic separation experiments: a) $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres 4 dispersed in water and b) after exposure to 1 T magnet for 1 min.

(0.24 wt%) in C% revealed that some sheets of rGO were removed from the surface of 4 during the stirring condition of reaction and resulted in the decrease of efficiency.

On the basis of band gap, a plausible mechanism was proposed for enhanced activity of $\text{rGO@CuZnO@Fe}_3\text{O}_4$ microspheres 4 for CO_2 reduction (Scheme 2). For the CO_2 reduction to methanol the position of conduction band of semiconductor should be more negative than reduction potential of $\text{CO}_2/\text{CH}_3\text{OH}$ (-0.38 V vs NHE at pH 7) and the position of valence band should be more positive than oxidation potential of $\text{H}_2\text{O}/\text{O}_2$ ($+0.82 \text{ V}$ vs NHE at pH 7) [3]. This means that the band gap of the semiconductor should be wide so both CO_2 reduction and water oxidation reaction can proceed simultaneously. Fe_3O_4 microspheres due to presence of Fe_2O_3 layer on the surface works as semiconductor and can produce electron hole pairs in visible light [42], but due to the narrow band gap (1.16 eV) they are unable to achieve water splitting. ZnO has a wide band gap (3.2 eV) with suitable positions of conduction band (-0.66 V vs NHE at pH 7) and valence band ($+2.54 \text{ V}$ vs NHE at pH 7) to reduce CO_2 to methanol and oxidize water to oxygen, respectively [55]. However, only UV light can make transition of electrons from valence to conduction band. So neither Fe_3O_4 nor ZnO alone could reduce CO_2 to methanol. However, when ZnO was coated on Fe_3O_4 the obtained microspheres can reduce CO_2 to methanol and achieve water splitting which can be explained on the basis of reduced band gap (1.90 eV) for $\text{CuZnO@Fe}_3\text{O}_4$ microspheres and synergistic mechanism of electron-hole pairs generation. After absorption of visible light, Fe_2O_3 present on the surface of Fe_3O_4 get excited and electron-hole pairs were produced in the conduction and valence band, respectively. Electrons from the conduction band of $\text{Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ move to the conduction band of ZnO while holes move to valence band of ZnO so indirectly electrons and holes were produced in CB and VB of ZnO, respectively. But due to electron-hole pair recombination the yield of methanol was found to be poor. *In-situ* addition of Cu slowed down the process of electron-hole pair recombination by capturing photogenerated electrons by the Cu present on the surface of the microspheres. GO wrapping further increases the photocatalytic performance of $\text{CuZnO@Fe}_3\text{O}_4$ because of efficient capturing of electrons by GO. After reduction GO to rGO on microspheres the combined band gap of matrix was found to be 1.44 eV. Wrapping of rGO not only reduces the band gap, but also increases photo efficiency to manifold by providing conductive surface so photogenerated electrons and holes move apart and can be used for photoreduction of adsorbed CO_2 to methanol. The



Scheme 2. Plausible mechanism of CO₂ reduction by rGO/CuZnO/Fe₃O₄ 4.

electrons and protons necessary for conversion of CO₂ to methanol were derived from water splitting by positive holes present in the valence band of ZnO. This is worthy to mention that the reduction potential of H⁺/H₂ is (−0.41 V vs NHE at pH=7) close enough to CO₂/CH₃OH (−0.38 V vs NHE at pH 7); hence we observed hydrogen as a byproduct along with the desired methanol [3].

4. Conclusions

We have developed novel magnetically separable core-shell structured reduced graphene oxide wrapped rGO/CuZnO/Fe₃O₄ microspheres that have been used as photocatalyst for the photoreduction of CO₂ to methanol under visible light irradiation. Owing to the synergistic effect of reduced graphene oxide and CuZnO/Fe₃O₄ in the rGO/CuZnO/Fe₃O₄ composite, significantly higher photocatalytic efficiency as compared to CuZnO/Fe₃O₄ as well as GO/CuZnO/Fe₃O₄ microspheres was observed. The methanol yield using CuZnO/Fe₃O₄, GO/CuZnO/Fe₃O₄ and rGO/CuZnO/Fe₃O₄ cat. was determined to be 858, 1749 and 2656 μmol g^{−1} cat, respectively. After completion of the reaction, the developed catalyst was easily recovered from the reaction mixture by an external magnet. The recovered catalyst showed consistent activity at least for six cycles and provided methanol in almost similar yields. As compared to the literature known photocatalysts, the developed photocatalyst offers several advantages such as high methanol yield, no need of sacrificial donor, facile recovery and efficient recycling.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.apcatb.2016.11.060>.

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Further reading

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